

WHAT IS CLAIMED IS:

- 1 1. A microanalysis chip comprising a body defining at least one
2 transfer-separation channel including a channel bottom having a bottom opening, the
3 transfer-separation channel terminating in a discharge aperture.
- 1 2. The microanalysis chip of claim 1 further comprising a seal
2 member positioned against a bottom surface of the body.
- 1 3. The microanalysis chip of claim 1 wherein the bottom opening
2 forms a well.
- 1 4. The microanalysis chip of claim 1 wherein the bottom opening
2 comprises a passive valve.
- 1 5. The microanalysis chip of claim 1 further comprising a reservoir in
2 the body and a reagent fluid in the reservoir.
- 1 6. The microanalysis chip of claim 1 comprising a plurality of the
2 transfer-separation channels.
- 1 7. The microanalysis chip of claim 1 wherein the bottom opening is
2 cooperatively structured to receive a pillar of a sample chip.
- 1 8. The microanalysis chip of claim 1 wherein the body comprises one
2 of silicon, glass, or polymeric materials.
- 1 9. The microanalysis chip of claim 1 further comprising a reservoir
2 and a reagent adapted to process proteins contained in the reservoir.
- 1 10. The microanalysis chip of claim 1 further comprising a fluid
2 distribution network.
- 1 11. The microanalysis chip of claim 1 further comprising a nozzle
2 containing the discharge aperture.
- 1 12. The microanalysis chip of claim 1 wherein the at least one
2 transfer-separation channel is positioned within the body.

1 13. The microanalysis chip of claim 1 further comprising a
2 chromatography/retention zone downstream of the bottom opening.

1 14. The microanalysis chip of claim 1 further comprising a lid and a
2 nozzle, wherein the lid has a nozzle.

1 15. A method for chemically affecting a sample comprising:
2 providing a microanalysis chip including a body having a
3 transfer-separation channel with a channel bottom having a bottom opening;
4 inserting a pillar into the bottom opening such that a sample supported by
5 the pillar communicates with the transfer-separation channel; and
6 passing a reagent fluid into the transfer-separation channel in order for the
7 reagent fluid to come in contact with the sample to chemically affect the sample.

1 16. The method of claim 15 wherein the pillar is on a base.

1 17. The method of claim 16 further comprising:
2 sealing a region between the microanalysis chip and the base with a seal
3 member.

1 18. A dispenser assembly comprising:
2 a dispenser chip including a dispenser body including a vertical channel;
3 and
4 a sample chip having a base and a sample structure, the sample structure
5 comprising a pillar and a sample surface, wherein the vertical channel of the dispenser
6 chip is cooperatively structured to receive the pillar.

1 19. The dispenser assembly of claim 19 further comprising:
2 a seal member between the dispenser body and the base of the sample
3 chip.

1 20. A microfluidic chip comprising:
2 a body having a bottom surface;
3 a plurality of discharge apertures; and
4 a plurality of transfer-separation channels in the body, each
5 transfer-separation channel defined by a channel bottom with a bottom opening, and

6 having a portion upstream of the bottom opening and a portion downstream of bottom
7 opening, and wherein each transfer-separation channel terminates at one of the discharge
8 apertures.

1 21. The microfluidic chip of claim 20 further comprising:
2 a plurality of reservoirs coupled to the transfer-separation channels.

1 22. The microfluidic chip of claim 20 further comprising:
2 a plurality of reservoirs; and
3 a plurality of delivery channels upstream of the plurality of
4 transfer-separation channels.

1 23. The microfluidic chip of claim 20 further comprising:
2 a plurality of nozzles, each nozzle containing one of the discharge
3 apertures.

1 24. The microfluidic chip of claim 20 further comprising:
2 a lid having a plurality of nozzles, each nozzle containing one of the
3 discharge apertures.

1 25. The microfluidic chip of claim 20 wherein the bottom opening
2 includes a passive valve.

1 26. The microfluidic chip of claim 20 wherein each transfer-separation
2 channel comprises a concentration/chromatography zone in the portion of the
3 transfer-separation channel downstream of the opening.

1 27. The microfluidic chip of claim 26 wherein the discharge apertures
2 are at a top surface of the microfluidic chip.

1 28. A microfluidic assembly comprising:
2 a microfluidic chip comprising (i) a body having a bottom surface, (ii) a
3 plurality of discharge apertures, and (iii) a plurality of transfer-separation channels in the
4 body, each transfer-separation channel defined by a channel bottom with a bottom
5 opening, and having a portion upstream of the bottom opening and a portion downstream
6 of bottom opening, and wherein each transfer-separation channel terminates at one of the
7 discharge apertures; and

8 a sample chip comprising a base including a non-sample surface and a
9 plurality of sample structures, each sample structure including a sample surface.

1 29. The microfluidic assembly of claim 28 wherein the sample surfaces
2 are elevated with respect to the non-sample surface.

1 30. The microfluidic assembly of claim 28 wherein each sample
2 structure comprises a pillar, wherein the sample surface is on the pillar.

1 31. The microfluidic assembly of claim 28 wherein the bottom opening
2 comprises a passive valve.

1 32. The microfluidic assembly of claim 28 further comprising:
2 a seal between the microfluidic chip and the sample chip.

1 33. The microfluidic assembly of claim 28 wherein the microfluidic
2 chip further comprises:
3 a plurality of reservoirs, each reservoir containing a reagent;
4 a plurality of delivery channels coupled to the plurality of reservoirs; and
5 a distribution network of fluid channels coupled to the plurality of delivery
6 channels.

1 34. A method of processing an analyte, the method comprising:
2 processing an analyte on a sample surface on an sample chip;
3 transferring the processed analyte through a transfer-separation
4 downstream of the sample surface, wherein the transfer-separation channel is in a
5 microfluidic chip above the sample chip; and
6 analyzing the processed analyte downstream of the sample surface.

1 35. The method of claim 34 wherein analyzing the processed sample
2 comprises analyzing the processed sample using mass spectrometry.

1 36. The method of claim 34 further comprising, prior to processing the
2 sample:
3 inserting the sample surface into a fluid channel in a dispenser chip,
4 wherein the sample surface is on a pillar;

5 depositing a liquid sample on the sample surface using the dispenser chip;
6 and
7 binding an analyte in the liquid sample to the sample surface.

1 37. The method of claim 34 wherein processing comprises:
2 dispensing a reagent on the sample surface; and
3 cleaving the analyte into subunits.

1 38. A microfluidic chip comprising:
2 a body having a bottom surface; and
3 a plurality of vertical channels in the body, wherein each opening is
4 cooperatively structured to receive a pillar of a sample chip.

1 39. The microfluidic chip of claim 38 wherein the body further
2 comprises:
3 a plurality of horizontal delivery channels in communication with the
4 plurality of vertical channels.

1 40. The microfluidic chip of claim 38 wherein the body further
2 comprises:
3 a plurality of reservoirs upstream of the plurality of vertical fluid channels.

1 41. The microfluidic chip of claim 38 the body comprises silicon,
2 glass, or polymeric materials.

1 42. The microfluidic chip of claim 38 wherein surfaces of the body
2 forming each vertical channel are hydrophobic.

1 43. The microfluidic chip of claim 38 wherein surfaces of the body
2 forming each vertical channel are hydrophilic.

1 44. A method of processing analytes, the method comprising:
2 inserting a plurality of sample surfaces into a plurality of vertical channels
3 in a dispenser chip, wherein the plurality of sample surfaces are on pillars of a sample
4 chip;
5 depositing a plurality of liquid samples on the sample surfaces while the
6 sample surfaces are in the vertical fluid channels;

7 binding analytes from the plurality of liquid samples to the sample
8 surfaces;
9 withdrawing the sample surfaces from the vertical fluid channels;
10 inserting the plurality of sample surfaces into a plurality of openings in a
11 microanalysis chip so that the plurality of sample surfaces are in communication with a
12 plurality of transfer-separation channels in the microanalysis chip; and
13 processing the analytes using reagents flowing through the
14 transfer-separation channels while the analytes are bound to the sample surfaces.

1 45. The method of claim 44 further comprising:
2 discharging the processed analytes from the microanalysis chip using a
3 plurality of nozzles in the microfluidic chip.

1 46. The method of claim 44 further comprising:
2 transferring the processed analytes to a mass spectrometer.

1 47. The method of claim 44 wherein the analytes are proteins, DNA, or
2 RNA.

1 48. The method of claim 44 wherein processing includes at least one of
2 derivatizing, cleaving, or unfolding the analyte.

1 49. The method of claim 44 wherein each vertical fluid channel
2 comprises a passive valve.

1 50. The method of claim 44 wherein each pillar has an aspect ratio
2 greater than about 1.

1 51. The method of claim 44 further comprising:
2 performing a chromatography process on the processed analytes.

1 52. The method of claim 44 further comprising:
2 separating the processed analytes from the sample surfaces; and
3 transferring the processed analytes downstream of the sample surfaces in
4 the transfer-separation channels.

1 53. A system for analyzing analytes, the system comprising:

2 an analysis assembly comprising (i) a microanalysis chip comprising a
3 body comprising at least one transfer-separation channel defined by a channel bottom
4 having a bottom opening, the transfer-separation channel terminating in a discharge
5 aperture, and (ii) a sample chip having a plurality of sample surfaces; and
6 an analysis device adapted to receive an analyte from the discharge
7 aperture.

1 54. The system of claim 53 wherein the analysis device is a mass
2 spectrometer.

1 55. The system of claim 53 wherein the sample surfaces are on pillars.